



Express Mail No.: EL 451 599 156 US

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

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Application of: Dasseux, *et al.*

Serial No.: 09/465,718

Filed: December 17, 1999

Group Art Unit: 1631

For: APOLIPOPROTEIN A-I  
AGONISTS AND THEIR USE  
TO TREAT DYSLIPIDEMIC  
DISORDERS

Examiner: Borin

Attorney Docket No.: 9196-018-999

Confirmation No.: 9219

**FEE TRANSMITTAL SHEET**

Assistant Commissioner for Patents  
Washington, D.C. 20231

Sir:

The fee required to be filed with the accompanying amendment of even date herewith concerning the above-identified application has been estimated to be \$0.

The claim amendment fee has been estimated as shown below:

(Col. 1)	(Col. 2)	(Col. 3)	SMALL ENTITY			OTHER THAN A SMALL ENTITY		
CLAIMS REMAINING AFTER AMENDMENT	HIGHEST NO PREVIOUSLY PAID FOR	PRESENT EXTRA	RATE	ADDIT. FEE	OR	RATE	ADDIT. FEE	
TOTAL 29	MINUS 62	= 0	× 9	\$ 0.00		× 18	\$	
INDEP. 01	MINUS 03	= 0	× 42	\$ 0.00		× 84	\$	
<input type="checkbox"/> FIRST PRESENTATION OF MULTIPLE DEP. CLAIM		140	\$	280				
			TOTAL	\$ 0.00	OR	TOTAL	\$	

Please charge the required fee to Pennie & Edmonds LLP Deposit Account No. 16-1150. A copy of this sheet is enclosed.

Respectfully submitted,



42,983

Date December 18, 2002

Rahul Pathak Reg. No.  
for Laura A. Coruzzi (Reg. No. 30,742)  
**PENNIE & EDMONDS LLP**  
1155 Avenue of the Americas  
New York, New York 10036-2711  
(650) 493-4935

Enclosure



12-27-02

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**AMENDMENT AND RESPONSE UNDER 37 C.F.R. § 1.114**

Commissioner for Patents  
Washington, D.C. 20231

Sir:

Reconsideration of the claims in light of the amendments and remarks that follow is kindly solicited. Enclosed herewith are Exhibits A (Claim Amendments: Version with Markings to Show Changes Made) and B (Claim Amendments: Pending Claims After Entry of the Instant Amendment), Fee Transmittal Sheet, Terminal Disclaimer, Terminal Disclaimer Fee Transmittal and Request for Continued Examination.

**AMENDMENT**

**TO THE CLAIMS**

Please cancel Claims 20-35, 43-55, 64-66 and 80-81 without prejudice.

Please amend Claims 1, 56-63, 67-75, 79, 82 and 83 to read as follows:

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1. (Twice amended) An ApoA-I agonist compound comprising:  
(i) a 15 to 26- residue peptide or peptide analogue according to formula (I) which forms an amphipathic  $\alpha$ -helix in the presence of lipids and exhibits at least about 38% LCAT activation activity as compared with human ApoA-I wherein one or two helical turns are